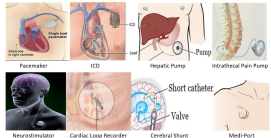

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Are You Ready When Patients Undergoing Radiotherapy Have Such Implanted Devices?

Maria Chan, PhD, DABR, DABMP, FAAPM
 AAPM Spring Clinical Meeting
 March 26-29, 2022


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This talk provides you with the essential information on managing commonly seen **implanted devices** for patients under radiotherapy




Identifying Implanted Devices (ID)

No relevant conflict of interest

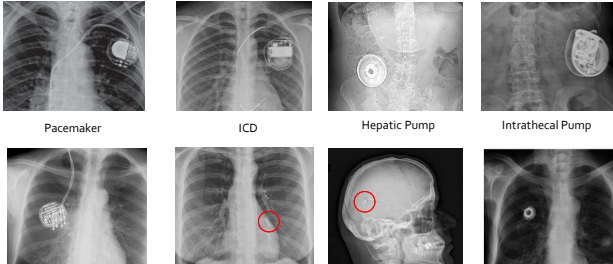


Categorizing & Managing ID



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Commonly seen implanted devices in Radiation Oncology

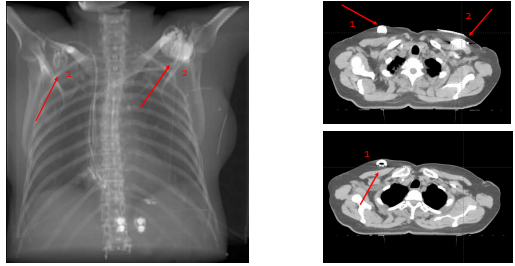


Pacemaker ICD Hepatic Pump Intrathecal Pump
 Neurostimulator Cardiac Loop Recorder Cerebral Shunt Medi-Port



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Picture credits: Wikipedia.org, Korean J of Ped, Willdoc, Dove Press, Radiopaedia, StatPearls Publishing, en.Wikipedia.org

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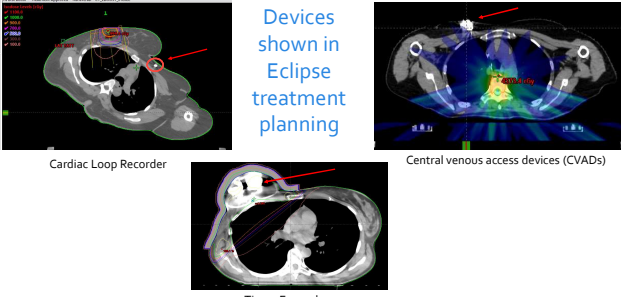


(1) Right Vascular Port; (2) Pacemaker



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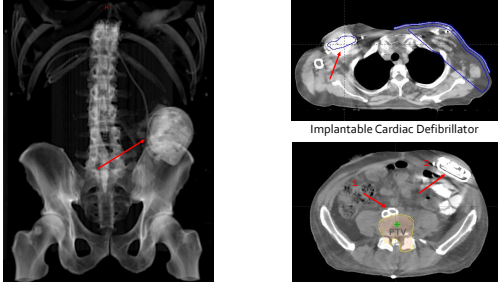
Devices shown in Eclipse treatment planning




Cardiac Loop Recorder Central venous access devices (CVADs)
 Tissue Expanders


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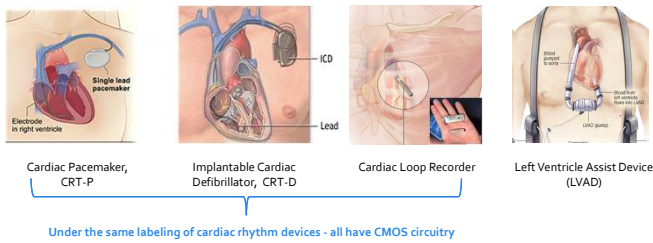


Intrathecal Pump in DRR Implantable Cardiac Defibrillator
 (1) Peripheral Artery Stent; (2) Infusion Drug Pump


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Patients with Cardiac Implantable Electronic Devices (CIEDs)

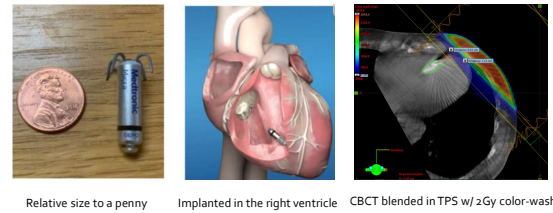


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Picture credit: NHLBI, Mayo Clinic

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Leadless Pacemaker (MR Compatible)

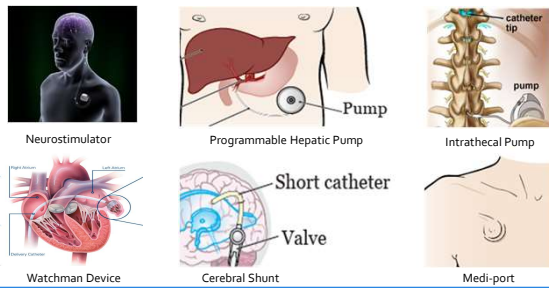


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Wang D, Chan MF, Zambri J, Lichtenwalner P, et al. *Advances in Radiation Oncology*. 2021. DOI:10.1016/j.adro.2021.100726

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Patients with Non-CIEDs Implanted Devices



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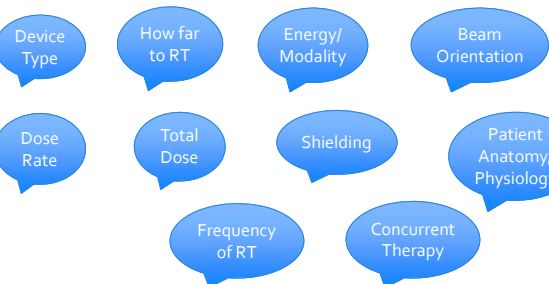
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Categorizing and Managing of Implanted Devices



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Factors Impacting Implanted Devices



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Implanted device	Clinical use	Susceptible component	Category	Dose limit	Historical malfunctions reported	References
Pacemaker	Control heartbeat	CMOS RAM Battery	Life-dependent	2-5 Gy	Transient damage Permanent damage EC damage Parameter reset Signal interference Battery depletion	8,9,18,19, 33,38,46
ICD	Sends electrical signals to the heart	CMOS RAM Battery	Life-dependent	0.5-2 Gy	Transient damage Permanent damage Pacing pulse change Pacing frequency change Sensing threshold change Lead impedance change Loss of telemetry capability Loss of signal/data Battery depletion	8,14,18,19, 33,38,39
Programmable hepatic pump	Gives continual chemo-therapy to the liver	EC, Battery	Adverse	10 Gy*	EC damage	20
Intrathecal pain pump	Gives continual pain medication to spine	EC, Battery	Adverse	26.5 Gy	Battery depletion Pump alarm	4,13
Neurostimulator	Sends electrical signals to the brain and spine	Implantable pulse generator	Adverse	5 Gy	NA	10,11,12
Loop recorder	Monitors heart rhythm	CMOS Battery	Adverse	5 Gy [†]	NA	5,39
Mediport	Vein access point for chemotherapy, IV, etc	NA	Without circuits	NA	NA	22
Cerebral shunt	Drains excess CSF from brain	NA	Without circuits	NA	NA	21

Abbreviations: CMOS = complementary metal oxide semiconductor; CSF = cerebrospinal fluid; EC = electronic circuit; IV = intravenous; NA = not applicable.

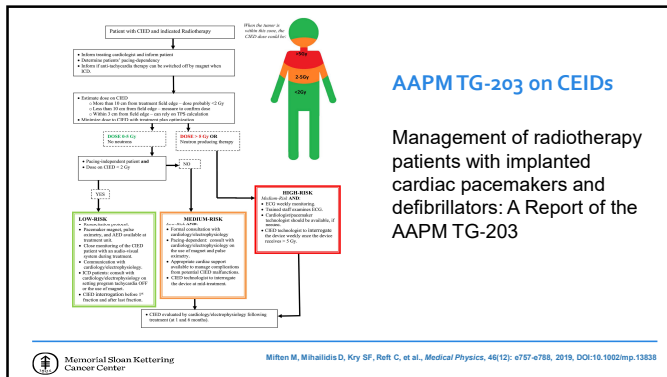
* Private communication from the manufacturer, Medtronic, Minneapolis, MN, October 23, 2019.

† A conservative model with the cardiologist on the safety of the information were able after radiation.

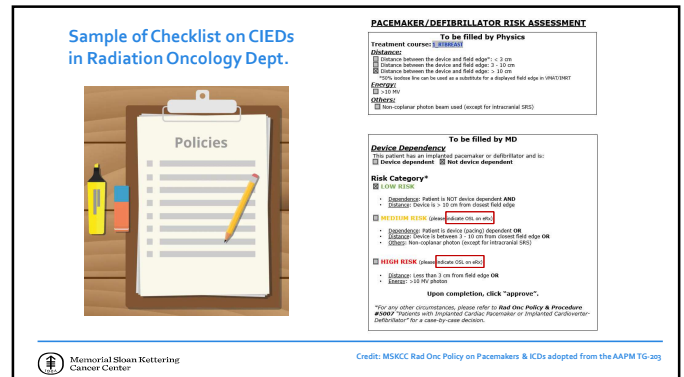
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Chan MF, Young C, Gelblum D, Shi C, et al. *Advances in Radiation Oncology*. 2021. DOI:10.1016/j.adro.2021.100732

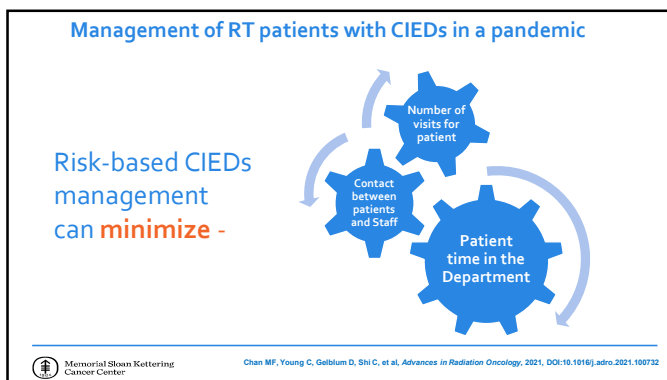
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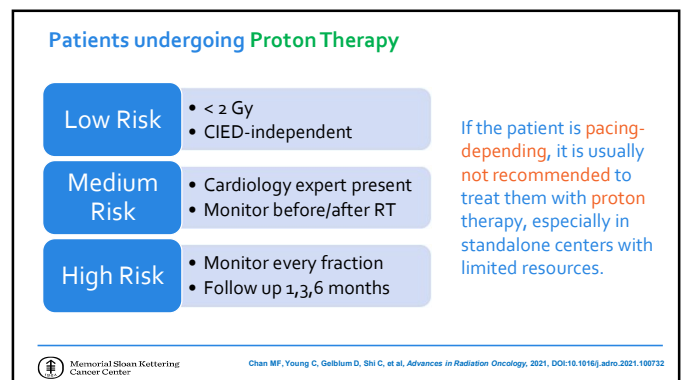
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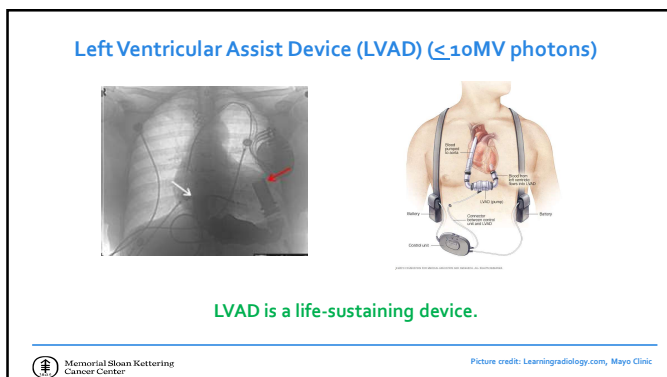
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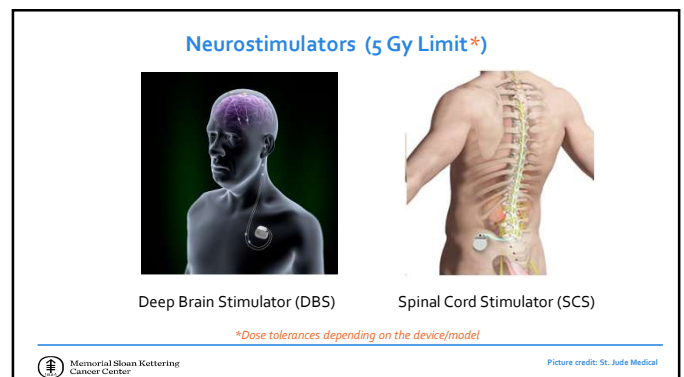
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Intrathecal Pump (28.5 Gy Limit*)

Pain relief medication

Intrathecal drug infusion system

Battery life 4-7 years

*Dose tolerances depending on the device/model

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Picture credit: Dove Medical Press

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Programmable Hepatic Pump (10 Gy Limit*)

10 Gy isodose color-wash display in Eclipse TPS

*Dose tolerances depending on the device/model

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Picture credit: MSKCC.org

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No Electronic Circuitry (No Dose Limit)

Brain Shunt

Medi-port

Central Venous Access Devices

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Picture credit: Wikipedia.org, Texas Winslow, StayWell

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Watchman Device™ (No Dose Limit)

Left atrial appendage closure (LAAC) therapy to prevent blood clots from forming and causing a stroke.

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Picture credit: Archgen

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A sample workflow to handle implanted devices

Treatment Planning

- Check IHS/EMR for implanted devices
- Risk assessment by MD
- Device Type: Cardiac Pacemaker, CRT-P, ICD, CRT-D; Programmable Hepatic Pump; Intrathecal Pain Pump; Neurostimulator; Loop Recorder***; Medi-Port; Cerebral Shunt
- High Risk*: Pacing dependent >1 Gy, or anatomy producing (e.g., >10 MV)
- Medium Risk*: <1 Gy and pacing independent, <1 Gy and pacing dependent
- Low Risk*: <1 Gy and pacing independent
- Discontinue dose limit and planning consideration (i.e., device known to reduce effect (due to device))
- Plan as normal, no special concern

Treatment Delivery

- CIED checked <24 h by CIED induction, ECG monitoring**
- Weekly check-up during RT; LIP study**
- Weekly check-up during RT; LIP study**
- ***Loop recorder - Cooperation needed with cardiologist on fidelity of information retrievable following radiation.
- Life-Dependency
- Adverse Effects
- Without Circuit

As new technology (i.e., ICD, CRT-D, etc.) evolves (1-2 copy on ICD of CIED -> copy on field edge)

**Recommended by Ref. (8); practices may vary

***Risk category based on AAPM TG-203, Ref. (18)

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In summary, proper policies on implanted devices should be made so you will be ready when your patients have such devices

- Life-dependent: Pacemaker, Implantable Cardiac Defibrillator, CRT-P, CRT-D, LVAD
- Adverse: Loop recorders, Neurostimulators, Programmable Hepatic Pumps, Intrathecal Pumps
- No electronic: Brain Shunt, CVAD, Medi-port, Watchman Device, Tissue Expander, Peripheral Artery Stent

Questions?

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